

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2003 ACS on STN

AN 2002:716267 CAPLUS

DN 137:247716

TI Preparation and use of substituted piperazine/piperidine derivatives as H receptor antagonists

IN Rosenblum, Stuart B.; Zeng, Qingbei; Mutahi, Mwangi Wa; Aslanian, Robert G.; Ting, Pauline C.; Shih, Neng-Yang; Solomon, Daniel M.; Cao, Jianhua; Vaccaro, Henry A.; McCormick, Kevin D.; Baldwin, John J.; Li, Ge

PA Schering Corporation, USA; Pharmacoepia, Inc.

SO PCT Int. Appl., 112 pp.

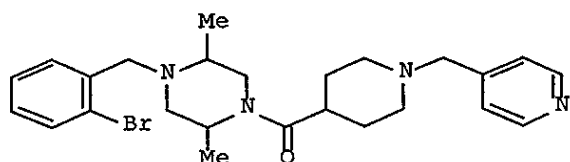
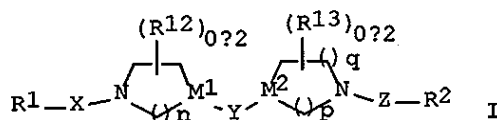
CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002072570	A2	20020919	WO 2002-US7106	20020311
	WO 2002072570	A3	20030306		
	W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, HR, HU, ID, IL, IN, IS, JP, KG, KR, KZ, LC, LK, LR, LT, LU, LV, MA, MD, MG, MK, MN, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UZ, VN, YU, ZA, ZM, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
	RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
	US 2003109564	A1	20030612	US 2002-95134	20020311
PRAI	US 2001-275417P	P	20010313		
OS	MARPAT 137:247716				
GI					



II

AB Title compds. I [R = (hetero)aryl, heterocycloalkyl, alkyl, carboxamido, etc.; X = alkyl, S(O)<sub>2</sub>; Y = bond, CO, CS, alkyl, amido, etc.; M = C, N; Z = alkyl, SO<sub>2</sub>, CO, carboxamido; R = 5-6 membered heteroaryl, alkyl, aryl, etc.; R = alkyl, OH, alkoxy, F, etc.; n, p, q = 1-3; with some provisions] were prepd. For instance, 2,5-dimethylpiperazine was alkylated with 2-bromobenzaldehyde (CH<sub>2</sub>Cl<sub>2</sub>, NaHB(OAc)<sub>3</sub>) and subsequently acylated with N-Boc-isonipecotic acid (CH<sub>2</sub>Cl<sub>2</sub>, PyBOP, i-Pr<sub>2</sub>NEt). The resulting intermediate was deprotected and reductively alkylated with pyridine-4-carboxaldehyde to afford. Selected example compds. had Ki within 0.2 and 600 nM for the H<sub>3</sub> receptor. : I, alone and in combination with a H<sub>1</sub> receptor antagonist, are used for the treatment of various diseases or conditions, such as, allergy, allergy-induced airway

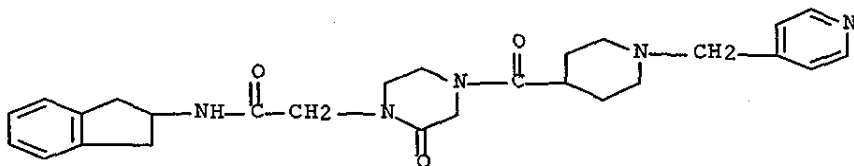
responses and congestion (e.g., nasal congestion).

IT 460093-95-2P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (H3 receptor antagonist; prepn. and use of substituted piperazine/piperidine derivs. as H receptor antagonists)

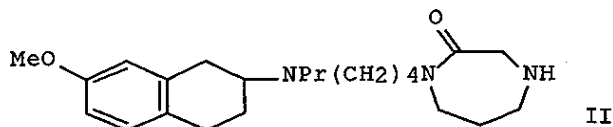
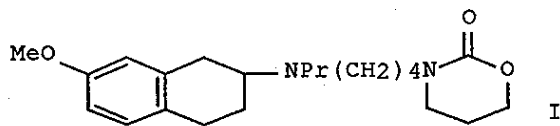
RN 460093-95-2 CAPLUS

CN 1-Piperazineacetamide, N-(2,3-dihydro-1H-inden-2-yl)-2-oxo-4-[[1-(4-pyridinylmethyl)-4-piperidinyl]carbonyl]- (9CI) (CA INDEX NAME)



L4 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2003 ACS on STN  
 AN 2001:868428 CAPLUS  
 DN 136:6017  
 TI Substituted 1-aminoalkyl-lactams and their use as muscarinic receptor antagonists  
 IN Madera, Ann Marie; Stabler, Russell Stephen; Weikert, Robert James  
 PA F. Hoffmann-La Roche A.-G., Switz.  
 SO PCT Int. Appl., '69 pp.  
 CODEN: PIXXD2  
 DT Patent  
 LA English  
 FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2001090082	A1	20011129	WO 2001-EP5631	20010517
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	EP 1289964	A1	20030312	EP 2001-933980	20010517
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
	BR 2001011019	A	20030617	BR 2001-11019	20010517
	JP 2003534331	T2	20031118	JP 2001-586271	20010517
	US 2002004494	A1	20020110	US 2001-862522	20010522
	US 6500822	B2	20021231		
	US 2003109524	A1	20030612	US 2002-289055	20021106
	US 6645958	B2	20031111		
	NO 2002005641	A	20021217	NO 2002-5641	20021122
PRAI	US 2000-207483P	P	20000525		
	US 2001-267617P	P	20010209		
	WO 2001-EP5631	W	20010517		
	US 2001-862522	A3	20010522		
OS	MARPAT 136:6017				
GI					



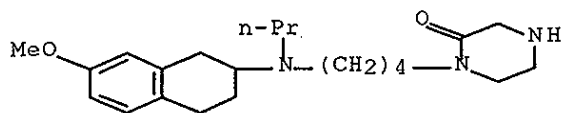
AB Title compds. such as I and II were prepd. Thus, I was prepd. in two steps from 3,4-dihydro-7-methoxy-2(1H)-naphthalenone and PrNH2. Muscarinic inhibitory activities (expressed as pKi values) of I were 8.20 (m2), 7.56 (m3), 6.30 (m5).  
 IT 375370-61-9P 375370-65-3P 375370-66-4P  
 375370-72-2P 375370-74-4P 375370-75-5P

375370-76-6P 375370-79-9P 375370-84-6P  
 375370-85-7P 375370-86-8P 375370-87-9P  
 375370-88-0P 375370-89-1P 375370-90-4P  
 375370-91-5P 375370-92-6P 375370-93-7P  
 375370-94-8P 375370-95-9P 375370-96-0P  
 375370-97-1P 375370-98-2P 375370-99-3P  
 375371-00-9P 375371-01-0P 375371-02-1P  
 375371-08-7P

RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)  
 (1-aminoalkyl-lactams and their use as muscarinic receptor antagonists)

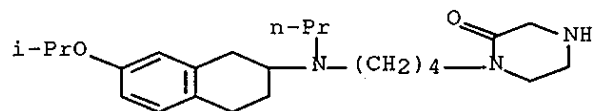
RN 375370-61-9 CAPLUS

CN Piperazinone, 1-[4-[propyl(1,2,3,4-tetrahydro-7-methoxy-2-naphthalenyl)amino]butyl]- (9CI) (CA INDEX NAME)



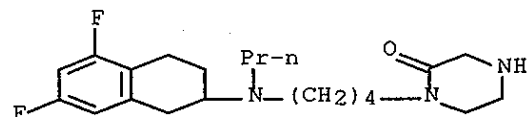
RN 375370-65-3 CAPLUS

CN Piperazinone, 1-[4-[propyl[1,2,3,4-tetrahydro-7-(1-methylethoxy)-2-naphthalenyl]amino]butyl]- (9CI) (CA INDEX NAME)



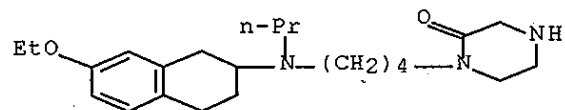
RN 375370-66-4 CAPLUS

CN Piperazinone, 1-[4-[(5,7-difluoro-1,2,3,4-tetrahydro-2-naphthalenyl)propylamino]butyl]- (9CI) (CA INDEX NAME)



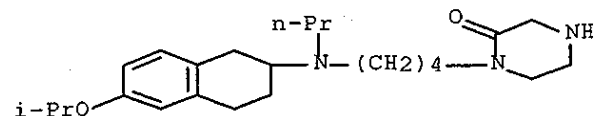
RN 375370-72-2 CAPLUS

CN Piperazinone, 1-[4-[(7-ethoxy-1,2,3,4-tetrahydro-2-naphthalenyl)propylamino]butyl]- (9CI) (CA INDEX NAME)



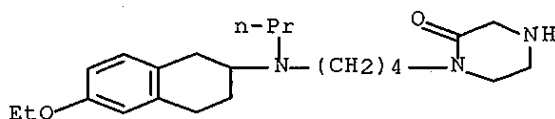
RN 375370-74-4 CAPLUS

CN Piperazinone, 1-[4-[propyl[1,2,3,4-tetrahydro-6-(1-methylethoxy)-2-naphthalenyl]amino]butyl]- (9CI) (CA INDEX NAME)



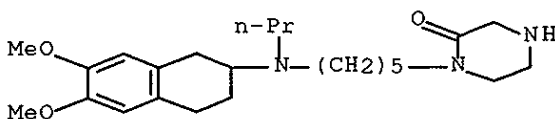
RN 375370-75-5 CAPLUS

CN Piperazinone, 1-[4-[(6-ethoxy-1,2,3,4-tetrahydro-2-naphthalenyl)propylamino]butyl]- (9CI) (CA INDEX NAME)



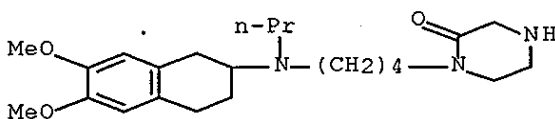
RN 375370-76-6 CAPLUS

CN Piperazinone, 1-[5-[propyl(1,2,3,4-tetrahydro-6,7-dimethoxy-2-naphthalenyl)amino]pentyl]- (9CI) (CA INDEX NAME)



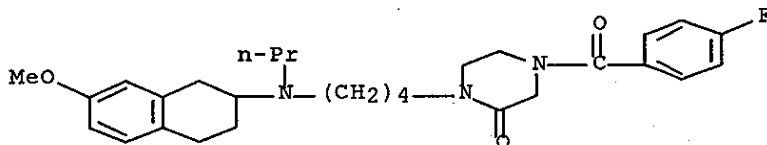
RN 375370-79-9 CAPLUS

CN Piperazinone, 1-[4-[propyl(1,2,3,4-tetrahydro-6,7-dimethoxy-2-naphthalenyl)amino]butyl]- (9CI) (CA INDEX NAME)



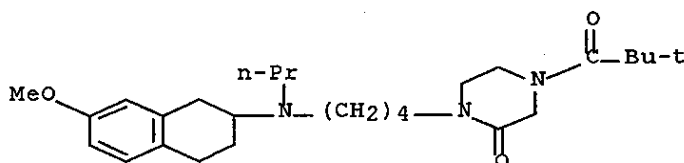
RN 375370-84-6 CAPLUS

CN Piperazinone, 4-(4-fluorobenzoyl)-1-[4-[propyl(1,2,3,4-tetrahydro-7-methoxy-2-naphthalenyl)amino]butyl]- (9CI) (CA INDEX NAME)



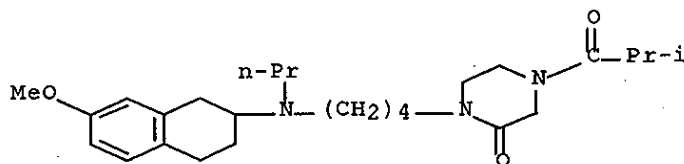
RN 375370-85-7 CAPLUS

CN Piperazinone, 4-(2,2-dimethyl-1-oxopropyl)-1-[4-[propyl(1,2,3,4-tetrahydro-7-methoxy-2-naphthalenyl)amino]butyl]- (9CI) (CA INDEX NAME)



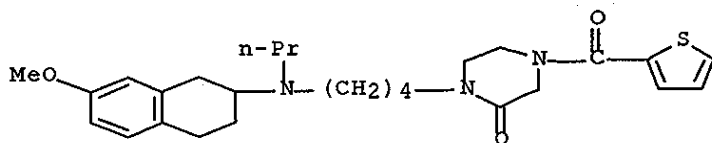
RN 375370-86-8 CAPLUS

CN Piperazinone, 4-(2-methyl-1-oxopropyl)-1-[4-[propyl(1,2,3,4-tetrahydro-7-methoxy-2-naphthalenyl)amino]butyl]- (9CI) (CA INDEX NAME)



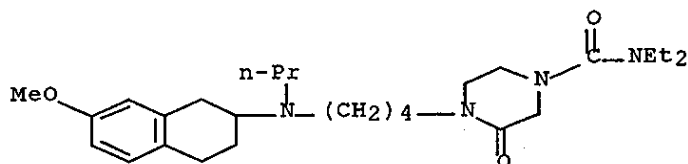
RN 375370-87-9 CAPLUS

CN Piperazinone, 1-[4-[propyl(1,2,3,4-tetrahydro-7-methoxy-2-naphthalenyl)amino]butyl]-4-(2-thienylcarbonyl)- (9CI) (CA INDEX NAME)



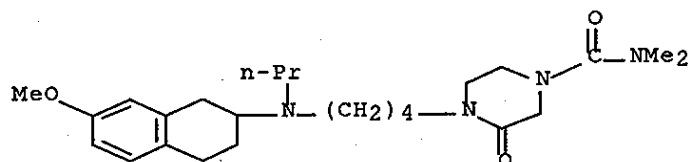
RN 375370-88-0 CAPLUS

CN 1-Piperazinecarboxamide, N,N-diethyl-3-oxo-4-[4-[propyl(1,2,3,4-tetrahydro-7-methoxy-2-naphthalenyl)amino]butyl]- (9CI) (CA INDEX NAME)



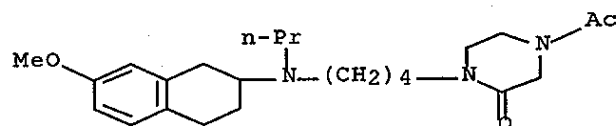
RN 375370-89-1 CAPLUS

CN 1-Piperazinecarboxamide, N,N-dimethyl-3-oxo-4-[4-[propyl(1,2,3,4-tetrahydro-7-methoxy-2-naphthalenyl)amino]butyl]- (9CI) (CA INDEX NAME)



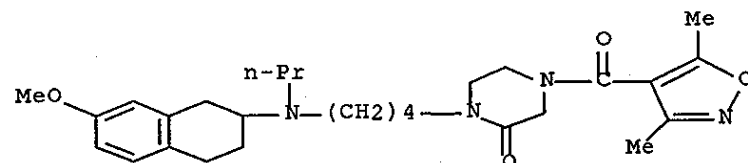
RN 375370-90-4 CAPLUS

CN Piperazinone, 4-acetyl-1-[4-[propyl(1,2,3,4-tetrahydro-7-methoxy-2-naphthalenyl)amino]butyl]- (9CI) (CA INDEX NAME)



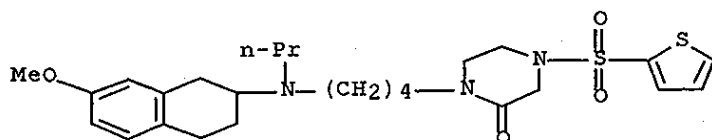
RN 375370-91-5 CAPLUS

CN Piperazinone, 4-[(3,5-dimethyl-4-isoxazolyl)carbonyl]-1-[4-[propyl(1,2,3,4-tetrahydro-7-methoxy-2-naphthalenyl)amino]butyl]- (9CI) (CA INDEX NAME)



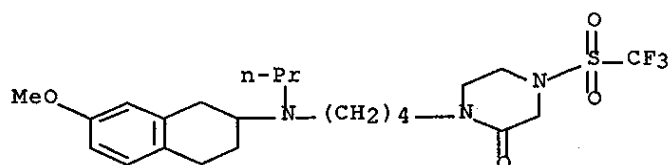
RN 375370-92-6 CAPLUS

CN Piperazinone, 1-[4-[propyl(1,2,3,4-tetrahydro-7-methoxy-2-naphthalenyl)amino]butyl]-4-(2-thienylsulfonyl)- (9CI) (CA INDEX NAME)



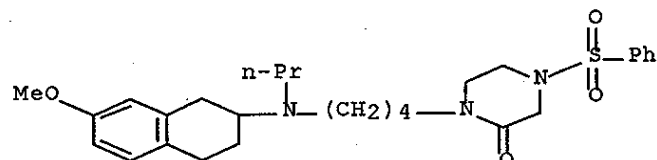
RN 375370-93-7 CAPLUS

CN Piperazinone, 1-[4-[propyl(1,2,3,4-tetrahydro-7-methoxy-2-naphthalenyl)amino]butyl]-4-[(trifluoromethyl)sulfonyl]- (9CI) (CA INDEX NAME)



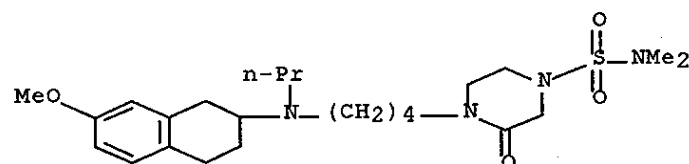
RN 375370-94-8 CAPLUS

CN Piperazinone, 4-(phenylsulfonyl)-1-[4-[propyl(1,2,3,4-tetrahydro-7-methoxy-2-naphthalenyl)amino]butyl]- (9CI) (CA INDEX NAME)



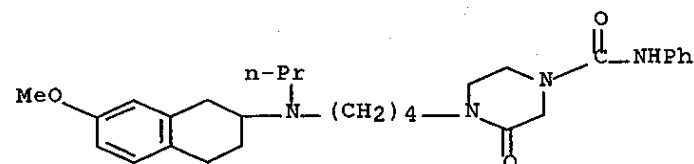
RN 375370-95-9 CAPLUS

CN 1-Piperazinesulfonamide, N,N-dimethyl-3-oxo-4-[4-[propyl(1,2,3,4-tetrahydro-7-methoxy-2-naphthalenyl)amino]butyl]- (9CI) (CA INDEX NAME)



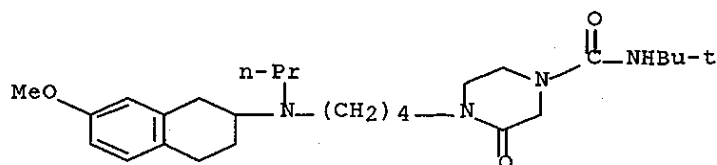
RN 375370-96-0 CAPLUS

CN 1-Piperazinecarboxamide, 3-oxo-N-phenyl-4-[4-[propyl(1,2,3,4-tetrahydro-7-methoxy-2-naphthalenyl)amino]butyl]- (9CI) (CA INDEX NAME)



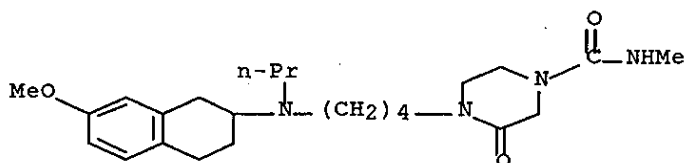
RN 375370-97-1 CAPLUS

CN 1-Piperazinecarboxamide, N-(1,1-dimethylethyl)-3-oxo-4-[4-[propyl(1,2,3,4-tetrahydro-7-methoxy-2-naphthalenyl)amino]butyl]- (9CI) (CA INDEX NAME)



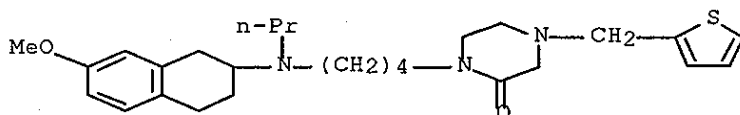
RN 375370-98-2 CAPLUS

CN 1-Piperazinecarboxamide, N-methyl-3-oxo-4-[4-[propyl(1,2,3,4-tetrahydro-7-methoxy-2-naphthalenyl)amino]butyl]- (9CI) (CA INDEX NAME)



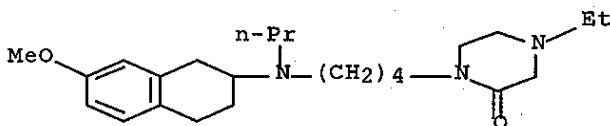
RN 375370-99-3 CAPLUS

CN Piperazinone, 1-[4-[propyl(1,2,3,4-tetrahydro-7-methoxy-2-naphthalenyl)amino]butyl]-4-(2-thienylmethyl)- (9CI) (CA INDEX NAME)



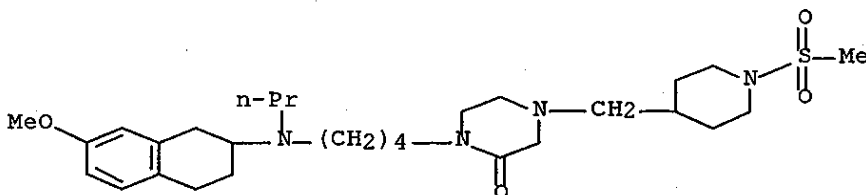
RN 375371-00-9 CAPLUS

CN Piperazinone, 4-ethyl-1-[4-[propyl(1,2,3,4-tetrahydro-7-methoxy-2-naphthalenyl)amino]butyl]- (9CI) (CA INDEX NAME)



RN 375371-01-0 CAPLUS

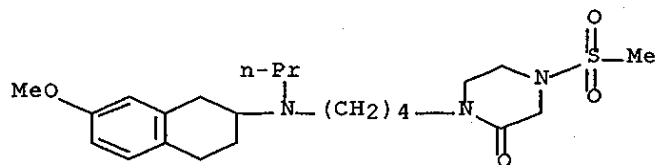
CN Piperidine, 1-(methylsulfonyl)-4-[[3-oxo-4-[4-[propyl(1,2,3,4-tetrahydro-7-methoxy-2-naphthalenyl)amino]butyl]-1-piperazinyl]methyl]- (9CI) (CA INDEX NAME)



RN 375371-02-1 CAPLUS

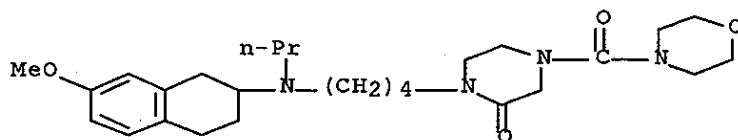
CN Piperazinone, 4-(methylsulfonyl)-1-[4-[propyl(1,2,3,4-tetrahydro-7-methoxy-2-naphthalenyl)amino]butyl]- (9CI) (CA INDEX NAME)





RN 375371-08-7 CAPLUS

CN Morpholine, 4-[[3-oxo-4-[4-[propyl(1,2,3,4-tetrahydro-7-methoxy-2-naphthalenyl)amino]butyl]-1-piperazinyl]carbonyl]- (9CI) (CA INDEX NAME)

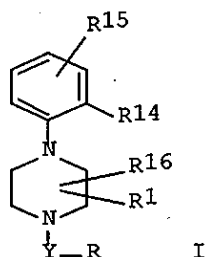


RE.CNT 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L10 ANSWER 1 OF 4 MARPAT COPYRIGHT 2003 ACS on STN  
 AN 129:27960 MARPAT  
 TI Preparation of piperazine derivatives as tocolytic oxytocin receptor antagonists  
 IN Bock, Mark G.; Evans, Ben E.; Culberson, J. Christopher; Gilbert, Kevin F.; Rittle, Kenneth E.; Williams, Peter D.  
 PA Merck and Co., Inc., USA  
 SO U.S., 37 pp., Cont.-in-part of U.S. 5,464,788.  
 CODEN: USXXAM  
 DT Patent  
 LA English  
 FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 5756504	A	19980526	US 1996-718415	19960923
	US 5464788	A	19951107	US 1994-217270	19940324
	WO 9525443	A1	19950928	WO 1995-US3738	19950323
	W: AM, AU, BB, BG, BR, BY, CA, CN, CZ, EE, FI, GE, HU, IS, JP, KG, KR, KZ, LK, LR, LT, LV, MD, MG, MN, MX, NO, NZ, PL, RO, RU, SG, SI, SK, TJ, TT, UA, US, UZ				
	RW: KE, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG				
PRAI	US 1994-217270		19940324		
	WO 1995-US3738		19950323		

GI

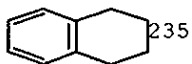


AB The title compds. I [Y = SO<sub>2</sub>, (CH<sub>2</sub>)<sub>p</sub>, CO(CH<sub>2</sub>)<sub>p</sub>, etc.; p = 1-3; R = (un)substituted Ph, etc.; R<sub>1</sub> = H, cyano, Ph, CONHR<sub>2</sub>, CONR<sub>2</sub>R<sub>2</sub>, etc.; R<sub>2</sub> = H, C3-8 cycloalkyl or C1-5 alkyl; R<sub>14</sub>, R<sub>15</sub> = C1-5 alkyl or alkoxy, halo; R<sub>16</sub> = H or oxo] were prepd. I are useful as oxytocin and vasopressin receptor antagonists. Thus, spiro[1H]indene-1,4'-piperidine.HCl was treated with 2,4-dimethoxy-phenylacetic acid in the presence of EDC, HBT and Et<sub>3</sub>N to give 1'-(2,4-dimethoxyphenylacetyl)-spiro[1H]indene-1,4'-piperidine, which showed IC<sub>50</sub> of 400 nM for [3H]oxytocin.

MSTR 2

~~181-4928-7~~<sup>5</sup>

G5 = 235

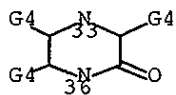


G28 = 128-161 129-71

~~128-129~~ G29

G29 = NH (SO)

G34 = 33-7 36-47



DER: or pharmaceutically acceptable salts

MPL: disclosure

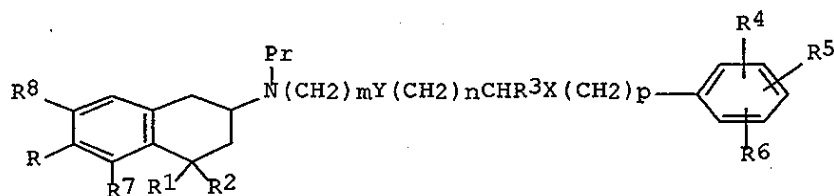
NTE: substitution is restricted

NTE: additional substitution also disclosed

RE.CNT 34 THERE ARE 34 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L10 ANSWER 2 OF 4 MARPAT COPYRIGHT 2003 ACS on STN  
 AN 125:86653 MARPAT  
 TI Preparation of 2-(N-propylamino)-1,2,3,4-tetrahydronaphthalene  
 dopaminergic D1 and D2 receptor agonist cardiovascular agents  
 IN Montanari, Stefania; Cavalleri, Paolo; Fraire, Cristina; Grancini, Gian  
 Carlo; Napoletano, Mauro; Santangelo, Francesco  
 PA Zambon Group S.P.A., Italy  
 SO PCT Int. Appl., 142 pp.  
 CODEN: PIXXD2  
 DT Patent  
 LA English  
 FAN.CNT 1

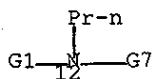
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PI	WO 9608228	A2	19960321	WO 1995-EP3562	19950911
	WO 9608228	A3	19960725		
	W:	AU, BY, CA, CZ, FI, HU, JP, KR, NO, NZ, PL, RU, SK, UA, US			
	RW:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG			
	US 5674909	A	19971007	US 1995-465636	19950606
	CA 2199484	AA	19960321	CA 1995-2199484	19950911
	AU 9535653	A1	19960329	AU 1995-35653	19950911
	AU 694563	B2	19980723		
	EP 781126	A2	19970702	EP 1995-932708	19950911
	EP 781126	B1	20011212		
	R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, NL, PT, SE			
	HU 76837	A2	19971128	HU 1997-1331	19950911
	JP 11501006	T2	19990126	JP 1995-509893	19950911
	RU 2149158	C1	20000520	RU 1997-106097	19950911
	AT 210433	E	20011215	AT 1995-932708	19950911
	ES 2168384	T3	20020616	ES 1995-932708	19950911
	FI 9701039	A	19970312	FI 1997-1039	19970312
	NO 9701134	A	19970512	NO 1997-1134	19970312
PRAI	IT 1994-MI1868		19940913		
	WO 1995-EP3562		19950911		
GI					



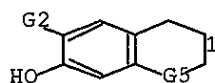
I

AB The title compds. (I; Markush definitions are provided within the document), useful for the treatment of arterial hypertension, congestive heart failure, renal failure, hypertension, and cerebrovascular insufficiencies, are prepd. Thus, (S)-N-propyl-N-[6-[(1,4-benzodioxan-2-yl)methylamino]hexyl]-5,6-dihydroxy-1,2,3,4-tetrahydro-2-naphthylamine dihydrochloride was prepd. and demonstrated a Ki of 0.66 nM against [3H]-domperidone on rat striated membrane-derived D2 receptors.

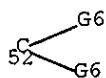
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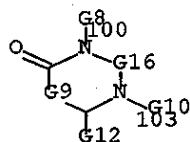
G1 = 1



G5 = 52



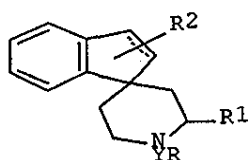
G8 = (4-8) CH2  
G16 = (1-2) CH2  
G21 = 100-12 103-14



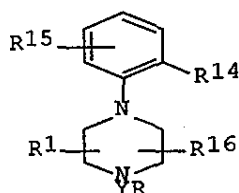
DER: and pharmaceutically acceptable salts  
MPL: claim 1  
NTE: substitution is restricted

L10 ANSWER 3 OF 4 MARPAT COPYRIGHT 2003 ACS on STN  
 AN 123:330860 MARPAT  
 TI Tocolytic oxytocin receptor antagonists  
 IN Bock, Mark G.; Evans, Ben E.; Culberson, J. Christopher; Gilbert, Kevin F.; Rittle, Kenneth E.; Williams, Peter D.  
 PA Merck and Co., Inc., USA  
 SO PCT Int. Appl., 114 pp.  
 CODEN: PIXXD2  
 DT Patent  
 LA English  
 FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9525443	A1	19950928	WO 1995-US3738	19950323
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	RW:	KE, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG			
	US 5464788	A	19951107	US 1994-217270	19940324
	CA 2186129	AA	19950928	CA 1995-2186129	19950323
	AU 9521952	A1	19951009	AU 1995-21952	19950323
	AU 686792	B2	19980212		
	EP 751773	A1	19970108	EP 1995-914875	19950323
	R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, NL, PT, SE			
	JP 09512521	T2	19971216	JP 1995-524838	19950323
	US 5756504	A	19980526	US 1996-718415	19960923
PRAI	US 1994-217270		19940324		
	WO 1995-US3738		19950323		
GI					



I

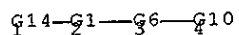


II

AB Spiroindenepiperidine derivs. I [R1 = H, C1-5 alkyl, CN, CO2H, Ph, etc.; R2 = H, PhCH2, C3-8 cycloalkyl, C1-5 alkyl; Y = CO2, C(O)NR2, C(:NR2), SO2, C(O)(CH2)n, (CH2)p, (CH2)pC(O); R = (tetrahydro)naphthyl, (substituted) cyclohexyl, (substituted) Ph, heterocyclyl; bond in cyclopentane ring is single or double; n = 0-3; p = 1-3] and phenylpiperazine derivs. II (Y, R, R1 as above; R14, R15 = H, C1-5 alkyl, C1-5 alkoxy, halo, NO2, CN; R16 = H, :O) and their pharmaceutically acceptable salts and esters are useful as oxytocin and vasopressin receptor antagonists for treatment of preterm labor and dysmenorrhea and for stopping labor prior to cesarean delivery. Thus, 1-[2-methoxy-4-[1-[2-(N-cyclopropylamino)ethylsulfonyl]-4-piperidyloxy]phenylacetyl]-4-(2-methylphenyl)piperazine-2-carboxamide (III) was prepd. in 11 steps from 4-hydroxypiperidine, Me 2,4-dihydroxybenzoate, 2-benzylaminoethanol, o-toluidine, 2,3-dibromopropionamide, and cyclopropylamine. III competed

with 1 nM oxytocin-3H for binding to rat uterine tissue with an IC50 of 20 nM.

**MSTR 1**

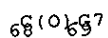


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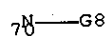


G5 = C(O)

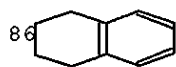
G6 = 68-2 69-4



G7 = 70



G10 = 86



DER: and pharmaceutically acceptable salts and esters  
MPL: claim 1  
NTE: substitution is restricted

L10 ANSWER 4 OF 4 MARPAT COPYRIGHT 2003 ACS on STN

AN 123:169654 MARPAT

TI Preparation of heterocyclic compounds as platelet aggregation inhibitors

IN Wayne, Michael Garth; Smithers, Michael James; Rayner, John Wall; Faull, Alan Wellington; Pearce, Robert James; Brewster, Andrew George; Shute, Richard Eden; Mills, Stuart Dennett; Caulkett, Peter William Rodney

PA Zeneca Ltd., UK

SO PCT Int. Appl., 236 pp.

CODEN: PIXXD2

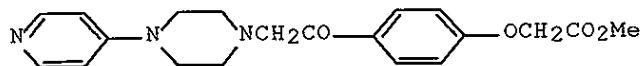
DT Patent

LA English

FAN.CNT 5

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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	WO 9422835	A3	19941222		
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	RW:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG			
	CA 2155307	AA	19941013	CA 1994-2155307	19940328
	AU 9462890	A1	19941024	AU 1994-62890	19940328
	AU 692439	B2	19980611		
	EP 690847	A1	19960110	EP 1994-910495	19940328
	R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT,			
SE	JP 08509967	T2	19961022	JP 1994-521811	19940328
	JP 3088016	B2	20000918		
	US 5750754	A	19980512	US 1996-658097	19960604
PRAI	GB 1993-6451		19930329		
	GB 1993-25610		19931215		
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	GB 1995-18188		19950907		

GI



I

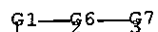
AB Title compds. [I; (M1)nQ(M2)1-nLA wherein = 0, 1; M1 = amino; Q = N-heterocyclyl; M2 = imino; L = template; A = an acidic group, or ester, amide deriv., sulfonamide] and pharmaceutically acceptable salts and pro-drugs thereof are prepd. Me 4-(bromoacetyl)phenoxyacetate in MeCN

was

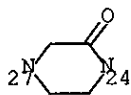
added to 1-(4-pyridyl)piperazine in MeCN to give the title compd II. Platelet aggregation inhibition was demonstrated by I. Pharmaceutical formulations comprising I are given.

MSTR 1A

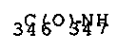




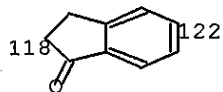
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G14 = 346-1 347-47

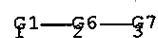


G15 = 118-46 122-3

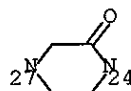


DER: or pharmaceutically acceptable salts or prodrugs  
 MPL: claim 1  
 NTE: additional ring formation is allowed  
 NTE: substitution is restricted

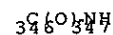
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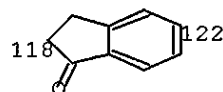
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G14 = 346-1 347-47

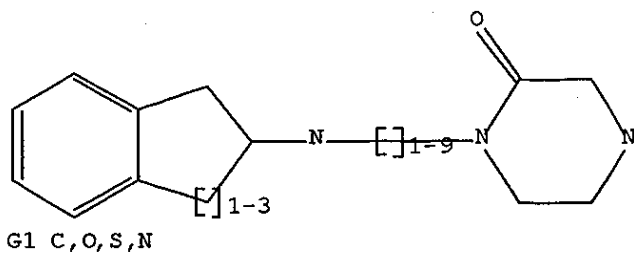


G34 = 118-46 122-48



DER: or pharmaceutically acceptable salts or prodrugs  
 MPL: claim 1  
 NTE: additional ring formation is allowed  
 NTE: substitution is restricted

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Structure attributes must be viewed using STN Express query preparation.

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 L3 29 S L1 FUL

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L4 2 S L3

FILE 'BEILSTEIN' ENTERED AT 18:56:12 ON 18 DEC 2003

L5 0 S L1  
 L6 0 S L1 FUL

FILE 'MARPAT' ENTERED AT 18:56:32 ON 18 DEC 2003

L7 0 S L1  
 L8 6 S L1 FUL  
 L9 5 S L8/COM  
 L10 4 S L9 NOT L4

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